STATUS OF THE CLAIMS

- 1.-16. (Cancelled).
- 17. (Currently Amended) A method of treating viral or bacterial infections, inflammatory diseases, or a tumor comprising tumor cells which express Hsp70 on their cell surface, said method comprising:
 - (a) administering a pharmaceutical composition consisting essentially of a pharmaceutically effective amount of isolated granzyme B as the only pharmaceutically active component to cells affected by said infection or inflammatory diseases or tumor cells, said tumor target cells bearing Hsp70 on their surface;
 - (b) allowing granzyme B to enter said cells via Hsp70 on the surface of said cells; and
 - (c) allowing said cells to undergo apoptosis as a result of the enzymatic activity of granzyme B.
- 18-24. (Cancelled).
- 25. (Previously Presented) The method of claim 17, wherein granzyme B is present in the pharmaceutical composition in a final concentration of 1µg/ml to 500µg/ml.
- 26. (Previously Amended) The method of claim 17, wherein granzyme B is present in the pharmaceutical composition in a final concentration of lng/ml to 10 ng/ml.
- 27. (Previously Amended) The method of claim 26 wherein granzyme B is present in the pharmaceutical composition in a final concentration of about 6 ng/ml.
- 28. (Cancelled).
- 29. (Currently Amended) A method of treating viral or bacterial infections, inflammatory diseases or a tumor comprising tumor cells which express Hsp70 on their cell surface, said method comprising:
 - (a) analyzing target tumor cells of a patient for surface expression of Hsp70;

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(b) administering a pharmaceutical composition comprising a pharmaceutically effective amount of granzyme B to cells affected by said infection or inflammation or said tumor cells, said cells bearing Hsp70 on their surface;

- (c) allowing granzyme B to enter said cells via Hsp70 on the cell surface; and
- (d) allowing said cells to undergo apoptosis as a result of the enzymatic activity of granzyme B.
- 30. (Previously Presented) The method of claim 29, wherein granzyme B is present in the pharmaceutical composition in a final concentration of 1μg/ml to 500μg/ml.
- 31. (Previously Presented) The method of claim 29, wherein granzyme B is present in the pharmaceutical composition in a final concentration of 1ng/ml to 10 ng/ml.
- 32. (Previously Presented) The method of claim 31 wherein granzyme B is present in the pharmaceutical composition in a final concentration of about 6 ng/ml.
- 33. (Cancelled)